Amendments to the Claims

Please amend the listing of claims as follows:

1. (Original) A compound of structural formula (I):

or a pharmaceutically acceptable salt or solvate thereof, wherein

Ar is:

aryl or heteroaryl which may both be substituted or unsubstituted;

 R_1 is independently:

hydrogen,

hydroxy,

cyano,

nitro,

halo,

alkyl,

alkoxy or

haloalkyl;

R₂ is:

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each R<sub>3</sub> is independently:
        hydrogen,
        halo,
        alkyl,
        haloalkyl,
        hydroxy,
        alkoxy,
        S-alkyl,
        SO<sub>2</sub>-alkyl,
        O-alkenyl,
        S-alkenyl,
        NR_7C(O)R_7
        NR<sub>7</sub>SO<sub>2</sub>R<sub>7</sub>,
        N(R_7)_2
        (D)-cycloalkyl,
        (D)-aryl,
        (D)-heteroaryl or
        (D)-heterocyclyl (wherein heterocyclyl excludes a heterocyclyl containing a
        single nitrogen), and
        wherein aryl, heteroaryl, heterocyclyl, alkyl and/or cycloalkyl may be
        substituted or unsubstituted, and two adjacent R<sub>3</sub> may form a 4- to
        7-membered ring;
R<sub>7</sub> and R<sub>8</sub> are each independently:
        hydrogen,
        alkyl or
        cycloalkyl, or
        R<sub>7</sub> and R<sub>8</sub> together with the nitrogen to which they are attached form a 5- to
        8-membered ring,
        wherein alkyl and cycloalkyl are both unsubstituted or substituted;
D is a bond or alkyl;
X is CH or N;
Y is O or NR<sub>7</sub>;
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n is 1 - 4;

m is 0 - 3;

o is 0 - 2;

p is 0 - 2;

q is 1 or 2;

s is 0 - 4.

2. (Original) The compound of claim 1, wherein

Ar is:

aryl which may be substituted with one to three substituents independently selected from the group consisting of cyano, nitro, perfluoroalkoxy, halo, alkyl, (D)-cycloalkyl, alkoxy and/or haloalkyl;

R₁ is independently:

hydrogen,

hydroxy,

halo,

alkyl,

alkoxy or

haloalkyl;

R₂ is:

$$R_8$$
 $(R_3)_{i}$

each R₃ is independently:

hydrogen,

halo,

alkyl,

haloalkyl,

hydroxy,

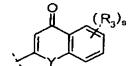
alkoxy,

S-alkyl or

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SO<sub>2</sub>-alkyl,
        O-alkenyl or
        S-alkenyl;
R<sub>7</sub> and R<sub>8</sub> are each independently:
        hydrogen,
        alkyl or
        cycloalkyl, or
        R<sub>7</sub> and R<sub>8</sub> together with the nitrogen to which they are attached form a 5- to
        7-membered ring optionally containing an additional heteroatom selected from
        O, S and NR<sub>4</sub>;
D is a bond or CH<sub>2</sub>;
X is CH or N;
Y is NR<sub>7</sub> or O;
n is 1 or 2;
m is 1 - 3;
o is 0 or 1;
p is 0 or 1;
q is 1;
s is 1 - 3.
(Currently Amended) The compound of claim 1-or 2, wherein
Ar is:
        phenyl or naphthyl which may be substituted with one or two substituents
        independently selected from the group consisting of perfluoroalkoxy, halo,
        alkyl, alkoxy and haloalkyl;
R<sub>1</sub> is independently:
        hydrogen,
        alkoxy,
        halo or
        alkyl;
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3.

R₂ is:



each R₃ is independently:

hydrogen,

hydroxy,

alkoxy,

SO₂-alkyl or

iso-propyl;

R₇ and R₈ are each independently:

hydrogen or

alkyl, or

R₇ and R₈ together with the nitrogen to which they are attached form a 6-membered ring optionally containing an additional oxygen atom;

X is CH or N;

Y is N-alkyl or O;

n is 1;

m is 1 - 3;

o is 0 or 1;

p is 0 or 1;

q is 1.

4. (Currently Amended) The compound of any of claims 1 to 3 claim 1, wherein Ar is:

phenyl or naphthyl which may be substituted with halo;

R₁ is hydrogen;

R₂ is:

each R₃ is independently:

hydrogen,

hydroxy,

alkoxy,

SO₂-alkyl or

iso-propyl;

R₇ and R₈ are each independently:

hydrogen or

alkyl, or

R₇ and R₈ together with the nitrogen to which they are attached form a 5- to 6-membered ring optionally containing an additional oxygen atom;

X is CH or N;

n is 1;

m is 1 or 2;

o is 0;

p is 0;

q is 1

s is 1 - 2.

5. (Currently Amended) The compound of any of claims 1 to 4 for use as a A medicament comprising the compound of claim 1.

- 6. (Currently Amended) Use of the compound of any of claims 1 to 4 for the preparation of a medicament for the treatment or prevention of A method of treating or preventing disorders, diseases or conditions responsive to the modulation of the melanocortin-4 receptor in a mammal, where modulation means activation in the case of MC4-R agonists or inactivation in the case of MC4-R antagonists, the method comprising administering to a human or mammal an effective amount of the compound of claim 1.
- 7. (Currently Amended) Use of MC4-R antagonists according to claims 6 for the preparation of a medicament for the treatment or prevention of cancer cachexia.

 A method of treating or preventing cancer cachexia, the method comprising administering to a human or mammal an effective amount of the MC4-R antagonists according to claim 6.
- 8. (Currently Amended) Use of MC4-R antagonists according to claims 6 for the preparation of a medicament for the treatment or prevention of muscle wasting.

 A method of treating or preventing muscle wasting, the method comprising administering to a human or mammal an effective amount of the MC4-R antagonists according to claim 6.
- 9. (Currently Amended) Use of MC4-R antagonists according to claims 6 for the preparation of a medicament for the treatment or prevention of anorexia. A method of treating or preventing anorexia, the method comprising administering to a human or mammal an effective amount of the MC4-R antagonists according to claim 6.
- 10. (Currently Amended) Use of MC4-R antagonists according to claims 6 for the preparation of a medicament for the treatment or prevention of anxiety and/or depression. A method of treating or preventing anxiety and/or depression, the method comprising administering to a human or mammal an effective amount of the MC4-R antagonists according to claim 6.

- 11. (Currently Amended) Use of MC4-R agonists according to claims 6 for the preparation of a medicament for the treatment or prevention of obesity. A method of treating or preventing obesity, the method comprising administering to a human or mammal an effective amount of the MC4-R antagonists according to claim 6.
- 12. (Currently Amended) Use of MC4-R agonists according to claims 6 for the preparation of a medicament for the treatment or prevention of diabetes mellitus.

 A method of treating or preventing diabetes mellitus, the method comprising administering to a human or mammal an effective amount of the MC4-R antagonists according to claim 6.
- 13. (Currently Amended) Use of MC4-R agonists according to claims 6 for the preparation of a medicament for the treatment or prevention of male or female sexual dysfunction. A method of treating or preventing male or female sexual dysfunction, the method comprising administering to a human or mammal an effective amount of the MC4-R antagonists according to claim 6.
- 14. (Currently Amended) Use of MC4 R agonists according to claims 6 for the preparation of a medicament for the treatment or prevention of erectile dysfunction.

 A method of treating or preventing erectile dysfunction, the method comprising administering to a human or mammal an effective amount of the MC4-R antagonists according to claim 6.
- 15. (Currently Amended) A pharmaceutical composition which comprises a compound of any of claims 1 to 4claim 1 and a pharmaceutically acceptable carrier.